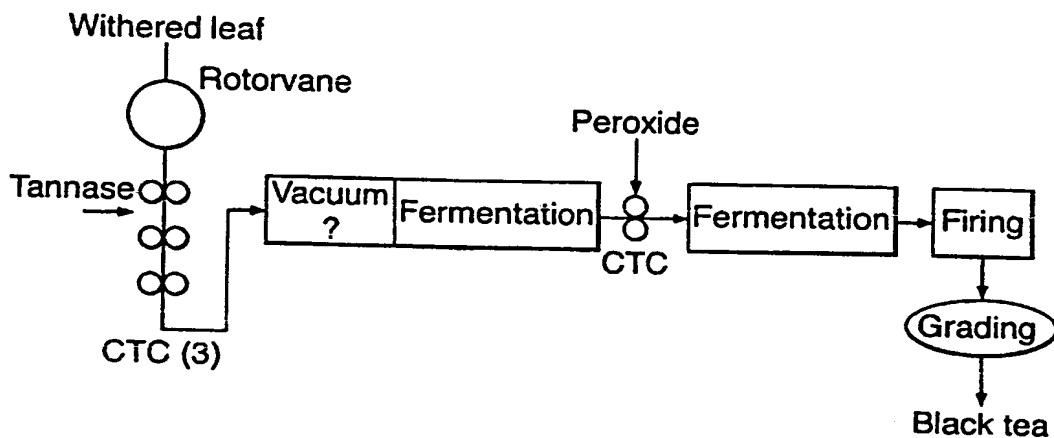


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## (57) Abstract

A method for preparing a cold water infusing leaf tea. Green tea leaves are macerated, treated with tannase, fermented in the presence of hydrogen peroxide in an amount that is sufficient to activate endogenous peroxidases to oxidise gallic acid and other compounds that are liberated by the tannase treatment, and then dried. The final product is a black leaf tea that infuses in hot or cold water to give good flavour and colour.

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## COLD WATER INFUSING LEAF TEA

5 The present invention relates to a method for manufacturing cold water infusing leaf tea. The method involves fermenting tannase pre-treated dhoor (macerated tea leaves) under solid-state conditions in the presence of hydrogen peroxide. The dried leaf product infuses in cold water to  
10 give good flavour and colour.

Background and prior art

15 Black leaf tea is traditionally produced by oxidising and drying freshly plucked green tea leaves. Tea, the beverage, is generally prepared in Commonwealth countries by brewing these tea leaves in freshly boiled water for a few minutes and adding milk, and perhaps a little sugar. However in  
20 some countries, notably the United States (or more accurately, parts thereof) tea is more commonly enjoyed as an iced beverage.

25 Such a beverage cannot be prepared conveniently by infusing traditionally manufactured tea leaves in cold water. Instead, Americans either infuse the leaves in hot water, remove the leaves and place the infusion in a refrigerator until it is ready to consume or place tea leaves in cold water in sunlight to infuse slowly over a period of hours.  
30

The numerous compounds in the leaves that give the beverage its unique organoleptical properties are only sparingly soluble in cold water. A more convenient option that has

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become available in the 1970's is the use of cold soluble tea-based powders.

5 There are numerous methods for making cold water soluble tea powders.

United States patent specification US 4,051,264 (Lipton/Sanderson) discloses a method for making a cold water soluble leaf tea extract. Tea leaves are pre-treated 10 with tannase under anaerobic conditions to generate a cold-water infusing tea with good colour, yield and flavour.

United States patent specification US 3,812,266 (Sanderson/Coggon) discloses a method that involves 15 converting green tea to black using tannase and natural tea enzymes. The method also includes a tannase pre-treatment step, but in a slurry system, followed by oxidation by natural tea enzymes to convert green tea into black, and generate tea powders, which are both hot and cold water 20 soluble. In some examples hydrogen peroxide is added, to "shorten the process". The proposed mechanism for enhanced cold-water soluble colour generation resulting from tannase-treatment (elevated epiteaflavic acid levels) is now known 25 to be incorrect, and no mechanism was presented to explain the effect of adding the hydrogen peroxide.

European patent specification EP 760,213 A1 (Unilever) discloses a method of enhancing colour in a tea-based foodstuff. The method involves using a tannase pre-treatment (on leaf or extract) followed by treatment with 30 exogenous peroxidase and hydrogen peroxide to generate cold-soluble colour.

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International patent publication WO 97/40699 (Unilever) concerns tea processing with zeolites to generate colour. There are examples of adding zeolite following tannase treatment to generate cold-water soluble tea.

5

United States patent specification US 4,639,375 (P&G, Tsai) discloses treating black tea with tannase, together with other cell-wall digesting enzymes, to generate cold-water soluble instant tea powders.

10

Convenient as cold water soluble tea powders can be, for many consumers the quality of the final beverage is not equal to that prepared from hot infused leaves. Other consumers prefer not to use powders as they perceive them to be artificial and therefore "unnatural".

The present inventors have surprisingly found that it is possible to make a leaf tea that infuses in cold water to give a beverage with good colour and flavour that is as acceptable to consumers as a hot infused black leaf tea that has been refrigerated. Furthermore this product can be made by modifying the traditional black tea manufacturing process.

Statement of the invention

In broad terms the present invention relates to a method for preparing a cold water infusing leaf tea comprising the  
5 steps of macerating green tea leaves, treating the macerated leaves with tannase, fermenting the tannase-treated macerate in the presence of an amount of hydrogen peroxide that is sufficient to activate endogenous peroxidases, and drying the fermented leaf material to yield the cold water  
10 infusible leaf tea.

The invention also relates to a method for generating colour species in a cold water soluble tea product comprising adding hydrogen peroxide to a tannase-treated macerate of  
15 green tea in a quantity that is sufficient for the endogenous peroxidases to oxidise gallic acid liberated by the tannase treatment.

"Tea" for the purposes of the present invention means leaf  
20 material from *Camellia sinensis* var. *sinensis* or *Camellia sinensis* var. *assamica*. It also includes rooibos tea obtained from *Aspalathus linearis* however that is a poor source of endogenous fermenting enzymes. "Tea" is also intended to include the product of blending two or more of  
25 any of these teas.

"Leaf tea" for the purposes of this invention means a tea product that contains one or more tea origins in an uninfused form.

30

"Cold water soluble" for the purposes of this invention means giving good colour, flavour and mouthfeel in a short

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infusion time i.e. less than 10 minutes, but preferably less than 5 minutes at a temperature at or above 4 °C.

The macerated leaves are preferably tannase treated under

5 anaerobic conditions. The process is effective without this anaerobic incubation provided sufficient tannase is used. The tannase treated macerate is preferably fermented under standard conditions to produce elevated levels of theaflavins and gallic acid prior to the addition of

10 hydrogen peroxide.

For the avoidance of doubt the word "comprising" is intended to mean including but not necessarily "consisting of" or "composed of". In other words the listed steps or options

15 need not be exhaustive.

Detailed description of the invention

20 Tea manufacture, especially black tea manufacture, traditionally comprises four basic steps: withering, rolling, fermenting and firing.

25 Withering is a process whereby the plucked tea leaves are stored for periods of time (perhaps up to 24 hours), during which they undergo various biochemical and physical changes which often includes a loss of moisture.

30 Maceration follows the withering step, and traditionally the withered leaves are optionally rolled to bruise and crush the leaves i.e. break down the plant tissue structure. This will have the effect of liberating fermentable substrates and fermenting enzymes from within the plant cells and

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tissue. Modern tea manufacture usually includes this step however the plant cells and tissue is broken down by passing tea, which has usually been withered, through a cutting machine.

5

The next step is commonly called fermentation but that is a misnomer. "Fermentation" is commonly used in the context of brewing alcohol to describe the action of exogenous enzymes. However in the tea world it is used to refer to the 10 oxidative process that tea undergoes when certain endogenous enzymes and substrates are brought together by mechanical disruption of the cells by tearing or cutting the leaves. During this process colourless catechins in the leaves are converted to a complex mixture of yellow and orange to dark- 15 brown substances and producing a large number of aromatic volatile compounds.

The colourful oxidation products include theaflavins and thearubigens. Theaflavins comprise several well-defined 20 catechin condensation products that are characterised by their benzotropolone ring. Thearubigens are a group of undefined molecules with a large variance in molecular weight. They have a large variety of colours ranging from yellow to dark red and brown.

25

The fermented product is fired and dried to give a black leaf tea. The firing involves heating and drying the tea to destroy the fermenting enzymes and thereby arrest 30 fermentation. It results in a reduction of moisture content to below 5%, and also leads to further chemical oxidation and changes in tea aroma. This generally involves exposing the tea to a blast of hot, dry air in a dryer.

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The present invention relates to method for making cold soluble black leaf tea. The present inventors have found that this can be achieved by modifying some of the steps of the traditional tea manufacturing process just described.

5

A preferred embodiment of the method of the invention is depicted in Figure 1. In that preferred embodiment freshly plucked green tea leaves are withered in the normal way using any of the art known techniques. Withering is not 10 essential to the invention but it can be a useful means to enhance tea aroma and also to reduce initial moisture content (which is important as moisture will be added with the tannase and peroxide, and drying efficiency can be affected at high moisture contents i.e. greater than 76%).

15

The leaves are macerated, which could mean being comminuted by a rotorvane and/or by a number of CTC (cut-tear-curl) machines.

20 In a first departure from traditional black tea manufacture the macerated leaves are treated with tannase (flavanol gallate esterase) to generate degallated catechins and gallic acid. This subsequently leads to the generation of theaflavins and non-gallated thearubigens during 25 fermentation (which are more soluble than the gallated ones).

30 The general reaction catalysed by tannase is the cleavage of gallate ester linkages, both on gallated catechins and also from other gallated compounds within the leaf. Tannase is also well known to improve the clarity of tea products since galloyl groups are important in cream formation and tannase

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has been used extensively for the degallation and solubilisation of black tea cream.

5 Tannase is known to be useful for treating green tea prior to slurry fermentation. For example the aforementioned US 3,812,266 (Sanderson et al) discloses using tannase to reduce the amount of tea cream in liquors. Improved colours generated by the process were also noted.

10 The present invention however does not require the tea to be suspended and fermented in a slurry. Indeed this would be counter-productive because the components necessary to give good colour and flavour from the leaf would be prematurely extracted into the slurry. Rather the tea is fermented

15 under solid-state conditions. This is an important distinction. The inventors previously thought that extraction of the catechins into a slurry liquor was essential for efficient action of tannase. They were surprised that direct application of tannase to the dhoool in

20 a solid-state led to efficient (i.e. almost complete) catechin degallation and high levels of theaflavin generation. They were even more surprised that the leaf product infused in cold water.

25 Tannase treatment degallates the gallated catechins ECG and EGCG to produce the degallated catechins EC and EGC. On subsequent oxidation during fermentation the catechins EC and EGC react to produce theaflavin.

30 The tannase can be applied using a variety of art-known techniques. The present inventors prefer to dissolve tannase in water, spray the solution onto the dhoool and leave the mixture to react for a suitable time at a suitable

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temperature. The tannase is applied to the dhoool after an initial maceration (for example, a first CTC cut) in a fine spray followed by subsequent cutting (for example, a second or third CTC cut) to ensure adequate mixing. The dhoool is 5 preferably incubated under vacuum, or under anaerobic conditions for example in an atmosphere or nitrogen. This prevents fermentation occurring. It is preferable that complete degallation takes place before fermentation starts as this results in maximal theaflavin levels in the 10 subsequent fermentation, which in turn leads to optimal colour generation.

The present inventors postulated that one might increase the efficiency by which certain exogenous compounds can access 15 certain endogenous compounds of solid state tea by using a vacuum to draw the exogenous molecules into the macerated tea leaves and into contact with the compounds therein. Vacuum infiltration per se is known. However it has been used to force substances between cells rather than into 20 cells. And those substances have tended to have small molecular weights.

The present inventors have however developed a method for bringing certain exogenous compounds into contact with 25 endogenous compounds found in tea that involves vacuum infiltrating macerated tea leaves with those exogenous compounds and applied it to modify certain properties of tea and tea based beverages. The extent to which this method allows even large molecules such as enzymes to access 30 endogenous tea compounds and modify certain properties of tea has been truly surprising. For example, an infusion of tannase pre-treated tea has been found to have more than

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double the total theaflavin content of a control and a six fold increase in TF1.

5 Vacuum infiltration is a technique that is often used in the preparation of protoplasts from plant tissue, albeit to introduce substances between rather than into cell walls. Cut leaf tissue is incubated in a solution containing tannase. The suspension is then placed under vacuum and air is drawn from the intracellular spaces of the leaf particles, the enzyme solution is drawn in to replace it. The inventors have found that a vacuum less than 100 mbar is suitable for this.

The major constraint when applying this method to tea dhoool  
15 is achieving access within the cells. Another major problem  
is that large volumes of water can seriously affect the  
quality of tea, by reducing oxygen uptake during  
fermentation. The results described in the Examples  
indicate that vacuum infiltration is a useful tool for  
20 introducing enzymes, for example tannase, into solid state  
dhoool. When fermented, tannase treated dhoool gives rise to  
black tea with high levels of theaflavin and no gallated  
species. This enables one to produce a range of novel teas.  
Vacuum assisted tannase treatment is much more effecgive in  
25 removing gallated species and enhacing additional TF  
formation than the equivalent treatment under ambient  
temperature and pressure. The vacuum allows the enzyme to  
penetrate the tissue and remove gallated species prior to  
fermentation, they key feature of tannase driven theaflavin  
30 enhancement, compared to simply applying the enzyme onto  
fermenting dhoool and mixing in by hand.

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If at all possible the conditions should be adjusted to prevent any fermentation prior to or during tannase treatment. This can be achieved by using a stronger vacuum pump, holding the dhool under N<sub>2</sub> sparge, or shortening the 5 tannase treatment.

10

Tannase can be applied to the macerated tea by a simple dosing. However, spraying the tannase in a fine mist is preferably as it aids infiltration.

Suitable conditions can be readily determined by experiment. Good results have been obtained with KIKKOMAN's tannase (KIKKOMAN is a trade mark of Kikkoman Corporation, Tokyo, Japan) in an amount of 1-100 mg/kg dhool, preferably 10-80 15 mg/kg dhool but more preferably 40-60 mg/kg dhool. Note: KIKKOMAN's tannase has 50,000 tannase activity units/gram.

20

Fermentation is preferably carried out at a pH in the range of 4.0 to 5.5. The fermentation temperature is preferably in the range 15 to 40 °C. Fermentation is preferably carried out for a time in the range 30 to 150 minutes, more preferably 105 to 120 minutes. However in a second departure from traditional black tea manufacture hydrogen peroxide is added, after a time that is sufficient to 25 generate gallic acid and theaflavin during the fermentation step, to activate (or at least greatly enhance the activity of) endogenous peroxidase.

30

Tea is known to contain natural peroxidase at high levels. It is also known that natural peroxidase can be activated (or have its activity enhanced) through the addition of hydrogen peroxide in a slurry system. J. Sci. Food Agric. vol. 32, p 920 - 932 (Dix., 1981) discloses such a system

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and process. The article mentions that peroxidase can oxidase tea polyphenols to form theaflavins and also thearubigens which may be similar and different to those produced under "normal" fermentations. However it does not 5 offer any detailed understanding as to the chemistry at work.

The present inventors have found that the endogenous peroxidases have the potential to oxidise catechins to 10 theaflavins and thearubigens, convert theaflavins to thearubigens and, unlike endogenous polyphenol oxidase, readily oxidise gallic acid. The combination of these reactions generates significant amounts of coloured 15 compounds that are soluble in cold water. The chemistry involved here is represented in Figure 2.

The hydrogen peroxide is added in an amount that is sufficient to activate endogenous peroxidases and oxidise gallic acid liberated by the tannase treatment. One skilled 20 in the art can determine that by experiment. However the present inventors prefer to use between 100 and 200 ml of 2.0 to 2.5% hydrogen peroxide per kg dhool, but preferably 160 ml of 2.0% hydrogen peroxide per kg dhool. Under normal conditions of tea manufacture peroxidase is largely 25 inactive, due to the low endogenous levels of hydrogen peroxide and high activities of catalase. Measurements have shown that all added hydrogen peroxide is consumed during the process, with none remaining in the final made tea. In contrast to the findings disclosed in the aforementioned US 30 4,051,264 the present inventors have found that the combination of tannase treatment and subsequent activation of peroxide is critical for the manufacture of a product

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that gives good colour and acceptable taste. Product that was only tannase treated had a "sour" or "metallic" note.

As one would expect, the colour and taste profile of a  
5 beverage made from the cold water infusing leaf tea of the present invention depends to a large extent on the source and quality of the raw material i.e. tea leaves. The present inventors have found that standard raw material, two leaves and a bud delivered to the factory within excess of  
10 1100 shoots per kilogram green leaf tea, can be processed according to the method of the invention to give very good colour and taste. However during efforts to improve the colour and taste even further the inventors surprisingly found significant improvement to both can be achieved by  
15 using more mature leaf.

Tea is generally harvested as two leaves and a bud on a 17 day cycle to optimise quality and yield. Extending the cycle means the leaves will be more mature and their  
20 chemical composition will be a little different. One then needs to pluck larger portions of tea plant to account for the extra growth. Such a harvesting strategy increases the yield per hectare of tea and thus improves productivity but the harvested plant material tends to have longer stems and  
25 a higher stem to leaf ratio.

Black tea made from mature tea leaf material tends to be less coloured and thinner after infusion than black tea made from tea portions of two leaves and a bud that are harvested  
30 on a 17 day cycle. However, the present inventors surprisingly found that when mature leaf tea, i.e. tea leaves harvested on a 30 to 50 day cycle in portions of 3 to 5, but preferably 4 leaves and a bud, is used there is a

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significant increase in cold water infusion performance and thus an improvement in colour and taste.

While not wanting to be bound by theory, it is thought that 5 mature leaf material (including stalk) contains higher levels of peroxidase than standard leaf and this peroxidase provides an important role in the maturation cascade. This means more peroxidase is available when hydrogen peroxide is added when carrying out the method of the invention.

10 Consequently more colour is generated by an enhanced peroxide/peroxidase oxidation system.

The method of the invention will now be described with reference to the following examples and the accompanying 15 drawings.

In the drawings:

Figure 1 is a diagram that shows a preferred process layout 20 of the invention.

Figure 2 is a diagram that represents the chemical reactions that tannase and endogenous peroxidase catalyse to give highly coloured thearubigens.

25 Figure 3 is a formulation profile of tannase - peroxide treated partially fermented dhool (Example 1).

Figure 4 is a histogram comparing the colour characteristics 30 of tannase-peroxide treated and untreated tea infusions (Example 3).

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Figures 5 and 6 represent the fermentation profiles for the experiments described in Example 4a.

Figures 7 and 8 show the composition of tannase-peroxide 5 processed black teas obtained using various tannase dosages (Example 5).

Figures 9a to 9e are infusion profiles of untreated and 10 tannase-peroxide treated teas at 4 °C, 15 °C, 25 °C, 55 °C and 70 °C respectively (Example 6).

#### **EXAMPLE 1**

15 **Lab scale process**

(a) **Process steps**

60 mg tannase dissolved in 24 ml water was sprayed onto 100 20 g of frozen dhool. The dhool was then thawed under N<sub>2</sub> and once it had reached 20 °C it was placed under vacuum (50 mbar) for 60 minutes. The dhool was then fermented for 60 minutes at 25 °C, 100% RH. After fermentation the dhool was sprayed with 12.5 ml ~2% hydrogen peroxide solution, placed 25 under vacuum for 15 minutes and then dried by a fluid bed drier (conventional FBD). One can increase the level of theaflavins by pre-treating the dhool with tannase. Peroxide is added to activate the endogenous tea peroxidase. This enzyme oxidises theaflavins and the gallic acid 30 released by tannase to give dark, cold water soluble pigments.

The fermentation profile is represented in Figure 3.

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(b) Taste testing

A cold water infusion made from tea prepared by the process just described was taste tested by an experienced tea 5 taster. The infusion was made in 200 ml of carbon filtered water for 5 minutes at 15 °C. The treated sample was described as having a "fruity note, good mouthfeel, astringent, lots of body, low aromatics, good colour, good ice tea product, stands up to ice" whilst the control, 10 standard black tea, was described as "bland". The results of colour, haze and solids analyses are given in Table 1 below where L\* is a measure of luminosity and a\* is a measure of red/green colour as determined using a MINOLTA (TM) colorimeter.

15

**Table 1: Lab scale process product**

	L*	a*	Haze	Solids
Control	54.6	23.6	26.8	0.2
Treated tea	29.2	34.6	29.4	0.24

20

**EXAMPLE 2**

Colour comparison with other products

25 Dhool was treated in treated in accordance with the basic lab scale process described above (60 mg tannase, dried immediately after peroxide addition). The colour of the liquor obtained from this product was compared with that 30 obtained from a commercially available product (SMOOTHBLEND

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(TM)) and a tea sample prepared from a standard fermentation of Kenyan tea leaf, all at 2.27 g black tea/200 ml chilled tap water, 5 minutes infusion. The results are summarised in Table 2 below. Note, these values are for tap water infusions which gives darker colours than infusions in distilled water.

10 TABLE 2 : Colour comparisons with other products

	L*	a*	b*
Tannase/peroxide	59.43	12.54	57.09
SMOOTHBLEND (TM)	94.22	-2.09	16.22
Kenyan	80.16	3.67	41.14

15 These results show the tannase/peroxide process on the invention gave a liquor whose colour was significantly brighter, redder and yellower than that obtained from the untreated Kenyan tea or the SMOOTHBLEND (TM) product.

20 EXAMPLE 3

Comparison of standard, tannase only, tannase peroxide (different times of addition) processes

25 Infusions were made for samples of a black tea standard, tannase treated black tea, tannase-peroxide treated black tea wherein the peroxide was added after 60 minutes fermentation, and tannase-peroxide treated black tea wherein the peroxide was added at  $t = 0$ . In each case 2.27 g tea  
30 was brewed for 5 minutes in 200 ml HIGHLAND SPRING (TM) still spring water at room temperature. The colour

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characteristics of each were measured using a MINOLTA transmission colorimeter. The results are represented in Figure 4.

5 These results and product tastings demonstrate that the combination of tannase treatment and hydrogen peroxide addition, after partial fermentation, are required to maximise the delivery of colour and flavour. Tannase only samples were observed to possess a "scur" or "metallic" 10 taste thought to be derived from the elevated levels of gallic acid.

The lab scale process described above was repeated and in addition comparison was made with an experiment where 15 tannase was not added, but in its place an amount of gallic acid equal to that released by tannase was added prior to hydrogen peroxide addition. The colour of the liquors obtained from the treated products were measured using a MINOLTA (TM) colorimeter. The results are given in Table 3 20 below where L\*, is a measure of luminosity, a\* is a measure of red/green colour, and b\* is a measure of yellow/blue colour.

25

TABLE 3 : Lab scale process variations

Treatment	controlled/infusion in distilled water	60 mg tannase + vacuum	Gallic acid - tannase
Infusion conditions	2.27g/200 ml. 5 min, 15 °C		
L*	90.33	84.65	91.63
a*	-1.33	2.27	-4.99
b*	47.36	80.43	52.57

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This experiment shows that addition of gallic acid alone is insufficient to generate cold-infusing colour/flavour.

5    **EXAMPLE 4**

Investigation of the use of a vacuum to optimise tannase effectiveness

10   The experiments that are described below were carried out using some standardised materials and methods. For completeness these are as follows:

Materials

15

Experiments were carried out with Kenyan leaf of BBK (Brooke Bond Kenya) clone 35, withered overnight at Mabroukie factory and air-freighted to the UK frozen on dry ice. This leaf was stored frozen at -80 °C until use. Tannase was from

20   Kikkoman's Co, Japan and "Macer8 W" (TM) from Biocatalysts Ltd, Wales.

Tannase treatment

25   Batches of dhool were prepared from frozen leaf of BBK 35 and stored at -80 °C. Portions of the batch were analysed for initial catechin composition and the remainder used in experiments. Experiments were carried out with various amounts of dhool; 8 g (for the standard experiment) to 100 g  
30   (for the scale up experiment). When more than 8 g was used the volume of tannase solution was scaled up appropriately.

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Eight grams of dhoor were thawed to room temperature under a stream of N<sub>2</sub>.

After tannase treatment some samples were fermented at 25 °C,  
5 100% RH, for up to 2 hours in a TEACRAFT (TM) controlled environment chamber. Tea was dried in a TEACRAFT(TM) fluid bed dryer at 120 °C air inlet temperature for 5 minutes then at 90 °C, for 20 minutes. At appropriate times portions of the dhoor were taken and immediately frozen in liquid  
10 nitrogen and stored at -80 °C until analysed.

In a variation upon this, a solution (1-2 ml) containing around 5 mg tannase (see results) was then pipetted onto the dhoor, trying to achieve an even spread. The flask was  
15 attached to a vacuum line for 45 to 60 minutes. Initially a bench top vacuum tap was used but in later experiments an EDWARDS (TM) vacuum pump was used as this provided a stronger vacuum.

20 Analysis of polyphenols

One gram of dhoor was refluxed for 30 minutes in 40 ml 70% (v/v) aqueous methanol. After cooling, the extracts were deleafed by filtering through 50 mm nylon mesh and their volume determined. A 200 µl aliquot was then added to 800  
25 µl antioxidant solvent and then analysed by reverse phase HPLC using diode array detection.

Dry weight measurement

The dry weight of dhoor samples were determined by mass  
30 difference following drying at 100 °C overnight. Results are expressed on a dry weight basis because moisture changes during N<sub>2</sub> sparge, and following tannase addition make a fresh weight basis unreliable.

- 21 -

Preparation of tea infusions

Infusions were prepared at 1% (w/v) tea solids, 5 minutes infusion time.

5

**EXAMPLE 4a**

Effectiveness of tannase against solid state dhool

10 In the first experiment, after thawing under N<sub>2</sub>, 8 g dhool was held under vacuum (using bench top vacuum tape) and 1 ml tannase solution was pipetted on to the dhool (5 mg/ml tannase, 31.25 µg (dhool)) and left for 60 minutes. Catechin composition was determined before and after this

15 treatment. The results are given in Table 4 below:

TABLE 4. Effect of tannase addition under Vacuum on catechin composition of BBK35 dhool

Moisture	µmol/g (DW)							
	G. acid	Caffeine	EGC	EC	EGCG	ECG	TF	
T=0 70.8	27	150	224	67	153	51	0	
T=60 77.4	227	137	303	107	5	0	13	
Change +6.6	+200	-13	+79	+40	-148	-51	+13	

20

Degallation was almost entirely complete with only 5 µmol/g (DW) residual EGCG after 60 minutes. The decrease in EGCG and ECG (total - 198 µmol/g (DW) was exactly mirrored by the increase in gallic acid (+200 µmol/g (DW)). However there was also evidence that some fermentation had taken place during the process; 13 µmol/g (DW) TF had formed and the increases in EGC and EC were not as high as predicted from

25

- 22 -

the decreases in EGCG and EC were not as high as predicted from the decreases in EGCG and ECG. There were no gallated theaflavins present. The increase in TF accounts for the missing EC/ECG but not the ECG/EGCG which suggests that some 5 thearubigins were formed as well. These results indicate that some fermentation was occurring during the tannase treatment and that perhaps small amounts of gallated theaflavins were being formed and then degallated. Although it was surprising that there was a degree of fermentation 10 under vacuum, it may be that the vacuum was not sufficiently strong to prevent it or that the apparatus used was not absolutely air tight. Nonetheless this proves that vacuum infiltration of tannase is very effective in degallating gallated catechins.

15

In the next experiment, the process was scaled up so that samples could be taken during the fermentation (to follow the profile of catechin oxidation) in solid state dhool following tannase treatment. Dhool (25 g) was thawed under 20 N<sub>2</sub> and treated with 15 mg tannase dissolved in 6 ml water and then held under vacuum for 60 minutes. At the end of this treatment the dhool was transferred to a controlled environment cabinet and fermented at 25 °C, 100% RH (relative humidity), for 2 hours. Samples were taken at 30 minute 25 intervals to follow the fermentation profile. The results are given in Table 5 below.

TABLE 5. Changes in catechins during Tannase treatment prior to fermentation

5

Once again degallation was almost complete with only 9  $\mu$ moles of EGCG remaining after 60 minutes in tannase treatment.

10 It would seem the increase in gallic acid accounted for the combined decrease in EGCG/ECG. However the increases in EGC and EC did not account for decreases in EGCG and ECG, which indicated that again some fermentation had taken place during the tannase treatment and indeed some theaflavin had formed. The fermentation profile for this experiment is shown in Figure 5 (see the drawings). EGC and EC were completely oxidised by 90 minutes, and theaflavin levels peak at over 60  $\mu\text{mol/g}$  DW (ie dry weight).

15

20 The enhancement in theaflavin following tannase pre-treated  
fermentation of solid state dhoor, is compared to a standard  
tea prepared from the same batch of dhoor in Figure 6 (see  
the drawings).

25 In any case these experiments demonstrate the vacuum infiltration enhances the ability of tannase to degallate catechin gallates and provides a raw material suitable for the formation of high levels of theaflavin during

- 24 -

fermentation. However, some fermentation is occurring during the tannase treatment.

5 EXAMPLE 4b

Preparation of black tea with enhanced theaflavin level

Having produced fermented dhoool with enhanced theaflavin 10 levels the next stage was to use this method to produce a tannase treated black tea so that the properties of the infusion could be assessed. The process was further scaled up to 100 g of dhoool so that the material could be fluid bed dried after fermentation. The dhoool (100 g) was thawed 15 under N<sub>2</sub> and 24 ml H<sub>2</sub>O containing 60 mg tannase added. The dhoool was then placed under vacuum, using an EDWARDS (TM) vacuum pump for 60 minutes. After this time the dhoool was placed in a controlled environment chamber and fermented at 25 °C, 100% RH for 120 minutes. After this the dhoool was 20 fluid bed dried. A replicate batch of tea was also prepared but without any tannase addition. Catechin levels before and after tannase treatment and after fermentation are shown in Table 6 below.

- 25 -

TABLE 6. Catechin and theaflavin  
Composition of tannase and control teas

	Mmol/g (DW)										
	G.a.	Caffeine	EGC	EC	EGCG	ECG	TF	TFMG	TF'MG	TfdiG	
t=0	32	192	231	72	205	78	0	0	0	0	
control	37	190	194	70	196	75	0	0	0	0	
t=60											
tannase	279	195	378	129	50	18	0	0	0	0	
t=60											
Net change	+246	+3	+14	+57	-155	-60	0	0	0	0	
T <sub>60</sub> -t <sub>0</sub>			7								
Control	23	189	0	10	8	5.5	2	2	2	3	
Fermented (t=180)	188	164	0	0	0	54	0	0	0	0	
Tannase Fermented (t=180)											

5

As one can see degallation of EGCG was only 75% complete, but there was good agreement between the reductions in EGCG and ECG and the increases in EGC and EC. Moreover there was 10 no theaflavin formation during the tannase treatment, this indicated that under stronger vacuum provided by the pump no fermentation was taking place.

Overall there is a 10 fold increase in TF1 and a 4.3 fold 15 increase in total TF was observed in the fermented dhoor (determined by solvent extraction).

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## EXAMPLE 5

Process optimisation5 (a) Optimisation of tannase dosage

Varying tannase dosage. A range of samples were prepared with increasing tannase dosage, between 5 and 320 mg tannase/kg. The results are given in Table 7 below where 10 L\*, a\* and b\* are as before.

15 TABLE 7 : Colour analysis of  
tannase/peroxide processed black teas

Tannase	L*	a*	b*	% soluble solids
5 mg/kg	87.27	0.77	60.9	0.215
10 mg/kg	86.62	1.42	62.9	0.205
20 mg/kg	86.54	0.56	64.7	0.210
40 mg/kg	86.3	1.6	68.1	0.225
80 mg/kg	82.3	5.62	77.5	0.230
160 mg/kg	81.8	6.4	82.9	0.245
320 mg/kg	81.7	6.5	83.7	0.260
Control	90.6	0.37	42.8	0.185

15 These results show the teas darkened (lower L-value) and  
20 increased in colour (a\* and b\*) with increasing tannase dosage. The inventors noted a step change in colour between using 40 and 80 mg tannase, with a 4 unit change in a\* and 5 unit change in b\*. Soluble solids increased with tannase dosage, maximum tannase gave 40% more soluble solids than  
25 the control.

- 27 -

The composition of tannase-peroxide processed black teas obtained from various tannase dosage are represented in Figures 7 and 8.

5 It is clear from these results that gallic acid, theaflavin and epitheafavic acid increase with tannase dosage, whilst residual catechins and gallated theaflavins fall.

10 (b) Further enhancements

Varying peroxide dosing

A comparison was made between adding water and different amounts of hydrogen peroxide, as above after 60 minutes 15 fermentation. To 1 kg of dhoool 80 mg tannase was added in 100 ml water, and after 60 minutes fermentation 160 ml of water or hydrogen peroxide solution (28 to 112 ml of 30% w/w was made up to 160 ml) was added. Colour measurements were made on the products and these are shown in Table 8 below.

20

TABLE 8: Peroxide dosing

Sample	L*	a*	b*	Solids
Water	87.39	1.61	78.53	0.26
HP1	83.08	6.22	88.38	0.27
HP2	82.57	5.85	86.39	0.22
HP3	81.87	5.79	80.38	0.22

HP1 - 28 ml peroxide + 132 ml water;

HP2 - 56 ml peroxide + 104 ml water;

25 HP3 - 112 ml peroxide + 48 ml water

- 28 -

There was little difference in colour between the different levels of hydrogen peroxide usage, but an indication that the level of solids extracted decreased with the higher amounts used.

5

Varying time of fermentation time after addition of peroxide

The length of fermentation after the addition of peroxide was varied from 0 to 60 minutes. There was little difference observed between 15 to 60 minutes, but addition 10 and immediate firing did result in a slight reduction in infusion colour.

Effect of CTC vs No CTC after peroxide application

Samples were prepared under standard conditions. Colour 15 measurements were made on the products and these are given in Table 9 below.

TABLE 9

Sample	L*	a*	b*	Solids
CTC	80.42	7.70	86.41	0.27
No CTC	81.65	7.42	87.81	0.26

20

This shows omitting the CTC after peroxide addition has a slight deleterious effect.

- 29 -

Effect of water volume during tannase application

Water volume was varied between 80 mg tannase in 60 to 160 ml water/kg dhoool. Colour measurements were made on the products and these are given in Table 10 below.

5

TABLE 10

Sample	L*	a*	b*	Solids
60 ml	83.25	3.90	81.49	0.24
80 ml	82.15	5.64	83.67	0.24
100 ml	81.98	5.68	85.75	0.25
120 ml	82.00	8.19	87.57	0.27
140 ml	82.21	5.53	80.97	0.22
160 ml	83.59	3.85	78.74	0.20

10 It was interpreted from these results that the optimum level for infusion performance 100 - 120 ml. The lower limit 100 was chosen in order to keep added moisture to a minimum.

Effect of increasing fermentation time

15 Fermentation time after tannase application was increased from 105 minutes (standard) to 120, 135, 150 minutes. Fermentation time after peroxide was kept at 15 minutes. The temperature was kept constant at 22 °F. Colour measurements were made on the products and these are given

20 in Table 11 below.

- 30 -

TABLE 11

Sample	L*	a*	b*	Solids
Std	84.12	4.24	86.10	0.22
120	83.25	5.18	90.03	0.20
135	83.22	5.08	89.21	0.21
150	81.64	6.93	92.60	0.22

These results indicated that infusion performance increased as fermentation time increased.

5

Effect of increasing fermentation temperature

10 Fermentation temperature was increased 22 °C, 25 °C, 30 °C, 35 °C, 40 °C. The longest fermentation time above was used (150 minutes + 15 minutes). Colour measurements were made on the products and these are shown in Table 12 below.

TABLE 12

Sample	L*	a*	b*	Solids
22C	81.64	6.93	92.60	0.22
25C	83.52	5.93	89.24	0.22
30C	81.59	6.33	80.89	0.22
35C	76.54	9.64	85.13	0.23
40C	79.20	7.09	82.95	0.22

15 It appeared from these results that performance increased as the temperature rose, up to a maximum of 30-35 °C was selected as optimum as this is the highest temperature consistently achievable during factory fermentation.

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Effect of multiple peroxide dosing (extended fermentation time and elevated temperature)

Standard peroxide (56 ml peroxide + 104 ml water), 15 minute ferment after addition. 2 Half peroxide (28 ml peroxide + 52 ml water), 10 minutes fermentation after each addition. 5 2 STD peroxide (56 ml peroxide + 24 ml water), 10 minutes fermentation after each addition. Colour measurements were made on the products and these are given in Table 13 below.

10

TABLE 13

Sample	L*	a*	b*	Solids
STD	80.33	5.29	83.77	0.24
2 1/2	83.46	3.10	79.54	0.21
2 STD	76.98	8.51	81.74	0.23

15 The results showed that doubling the dose of peroxide gave a benefit but splitting the standard dose into two applications seemed to reduce the infusion performance.

**EXAMPLE 6**

20 Infusion rate comparison

The infusion rate of 1.0 g for untreated tea and tannase-peroxide tea in 300 ml was determined by measuring the absorbance at 445 nm at various time intervals. These tests

were conducted at 4 °C, 15 °C, 25 °C, 55 °C and 70 °C respectively. The results are given in Figures 7a to 7e (Note C.I. in the drawings means "cold infusing" i.e. the tannase-peroxide treated tea). As one can see, the tannase-peroxide treated tea infused faster and to a greater extent at all temperatures.

**EXAMPLE 7**

10

### Mature leaf tea processing

15 Mature leaf tea, tea leaf material harvested in a 35 day cycle in portions of four leaves and a bud, was used as the raw material for the lab scale process described in Example 1. The infusion performance of the resulting cold water infusing leaf was measured as before using a MINOLTA (TM) colorimeter. The results are given in Table 14 below.

20

Table 14 - infusion performance of product made from mature tea leaf material

	L*	a*	B*
Run 1	71.4	13.4	85.5
Run 2	72.1	12.1	84.2
Run 3	78.2	5.7	75.7
Run 4	75.7	7.9	73.9
Run 5	74.2	6.8	78.4
Standard Leaf	81.8	3.2	74.5

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When compared with the typical infusion performance of products made from standard leaf about a 10 L-unit gain in colour was seen, decreasing to 3 L-units in some cases. In all cases the taste profile of the samples made using mature tea leaf material were preferred over the control.

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## CLAIMS

1. A method for preparing a cold water infusing leaf tea comprising the steps of macerating green tea leaves, 5 treating the macerated leaves with tannase, fermenting the tannase-treated macerate for a time that is sufficient to generate gallic acid and theaflavin, continuing the fermentation in the presence of an amount of hydrogen peroxide that is sufficient to activate endogenous 10 peroxidases, and drying the fermented leaf material to yield the cold water infusible leaf tea.
2. A method according to claim 1, wherein the macerated leaves are tannase treated under anaerobic conditions. 15
3. A method according to claim 2, wherein the tannase is added in a concentration between 10 and 80 mg/kg dhool.
4. A method according to any preceding claim, wherein the 20 macerated tea leaves are vacuum impregnated with the tannase.
5. A method according to any preceding claim, wherein the hydrogen peroxide is added in a concentration of between 100 25 and 200 ml of 2.0 to 2.5% hydrogen peroxide per kg dhool.
6. A method according to claim 5, wherein the hydrogen peroxide is added in a concentration of about 160 ml 2% peroxide per kg dhool. 30
7. A method according to claim 5 or 6, wherein the hydrogen peroxide is added after a time that is sufficient

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to generate gallic acid and theaflavin during the fermentation and thereby activate endogenous peroxidase.

8. A method according to any preceding claim, wherein the  
5 green tea leaves include mature tea leaf tea material.

9. A method according to any preceding claim, wherein said  
mature tea leaf material comprises tea portions of 3 to 5  
leaves and a bud harvested in a 30 to 50 day cycle.

10

10. A method for generating colour species in a cold water  
infusing tea product comprising adding hydrogen peroxide to  
a preferably fermented tannase-treated macerate of green tea  
in an amount that is sufficient for the endogenous  
15 peroxidases to oxidise gallic acid liberated by the tannase  
treatment and oxidise the theaflavin produced.

20

11. A cold water infusing leaf tea obtainable by macerating  
green tea leaves, treating the macerated leaves with  
tannase, fermenting the tannase-treated macerate for a time  
that is sufficient to generate gallic acid and theaflavin,  
continue the fermentation in the presence of an amount of  
hydrogen peroxide that is sufficient to activate endogenous  
peroxidases, and drying the fermented leaf material to yield  
25 the cold water infusing leaf tea.

30

12. A cold water infusing leaf tea according to claim 10,  
wherein the green tea leaves include mature tea leaf tea  
material.

13. A cold water infusing tea powder made from the cold  
water infusing leaf of claim 10 or 11.

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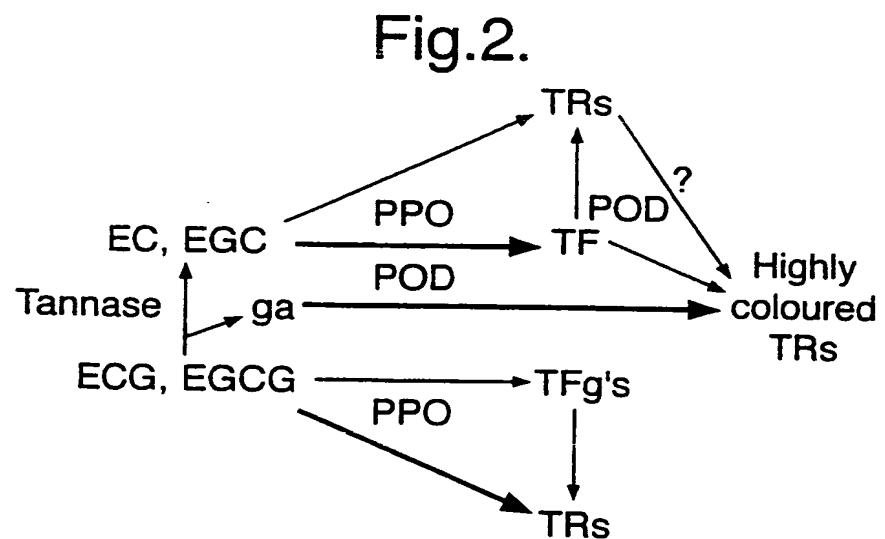
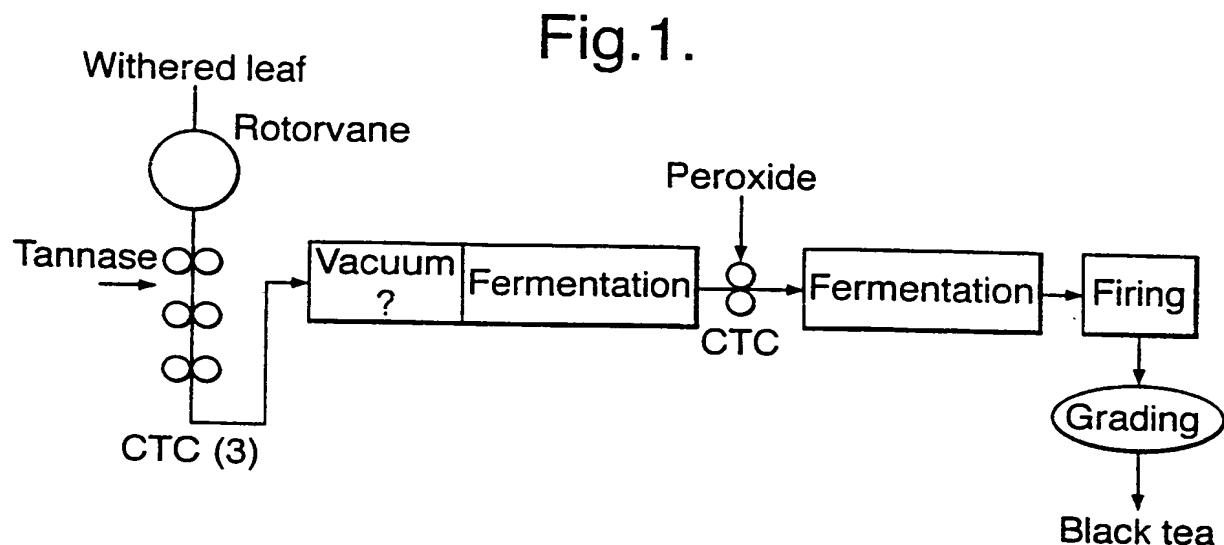


Fig.3.

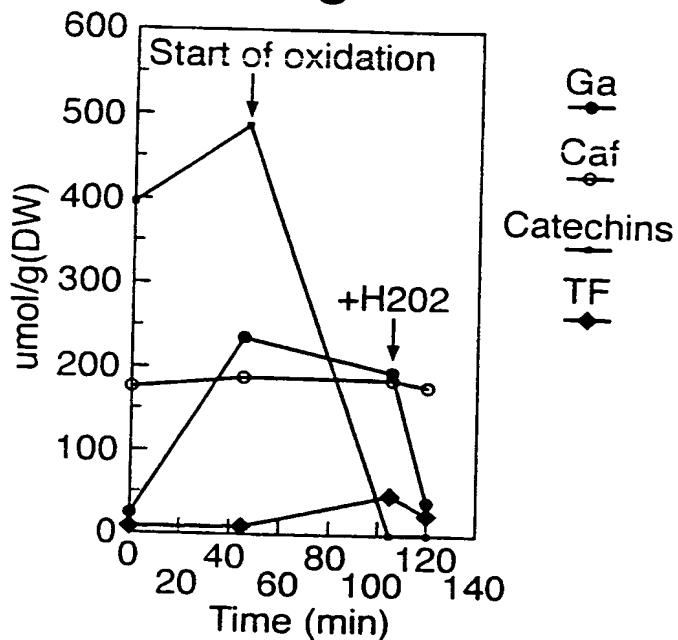


Fig.4.

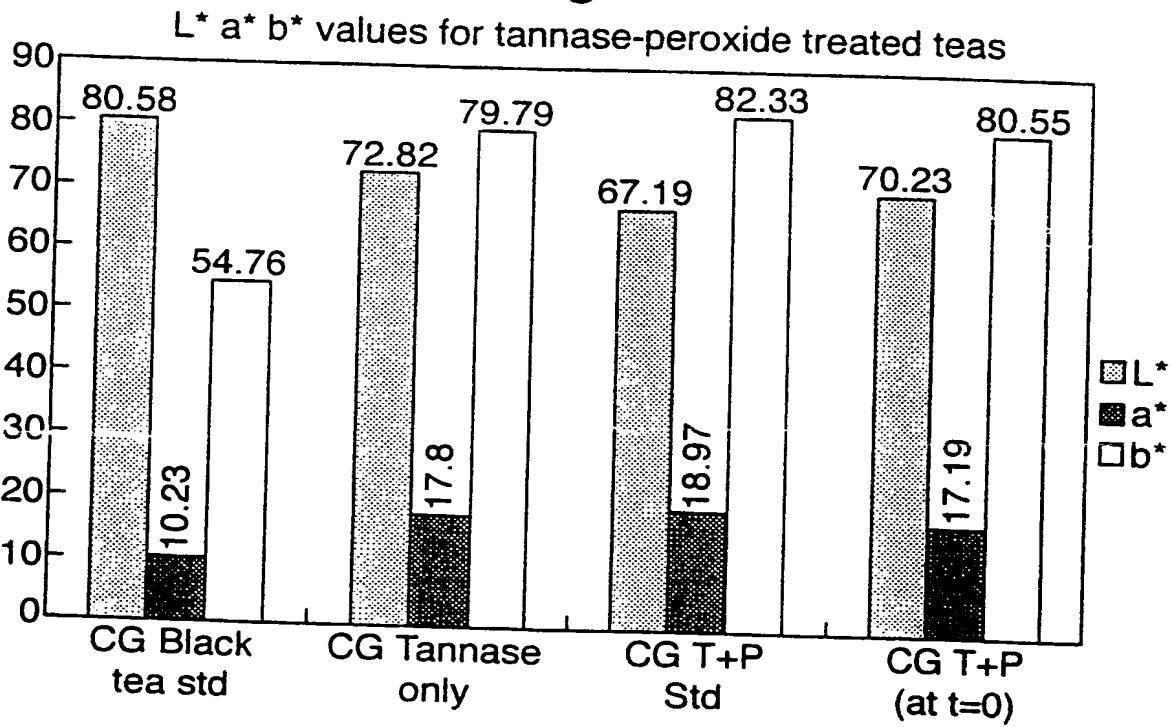


Fig.5.

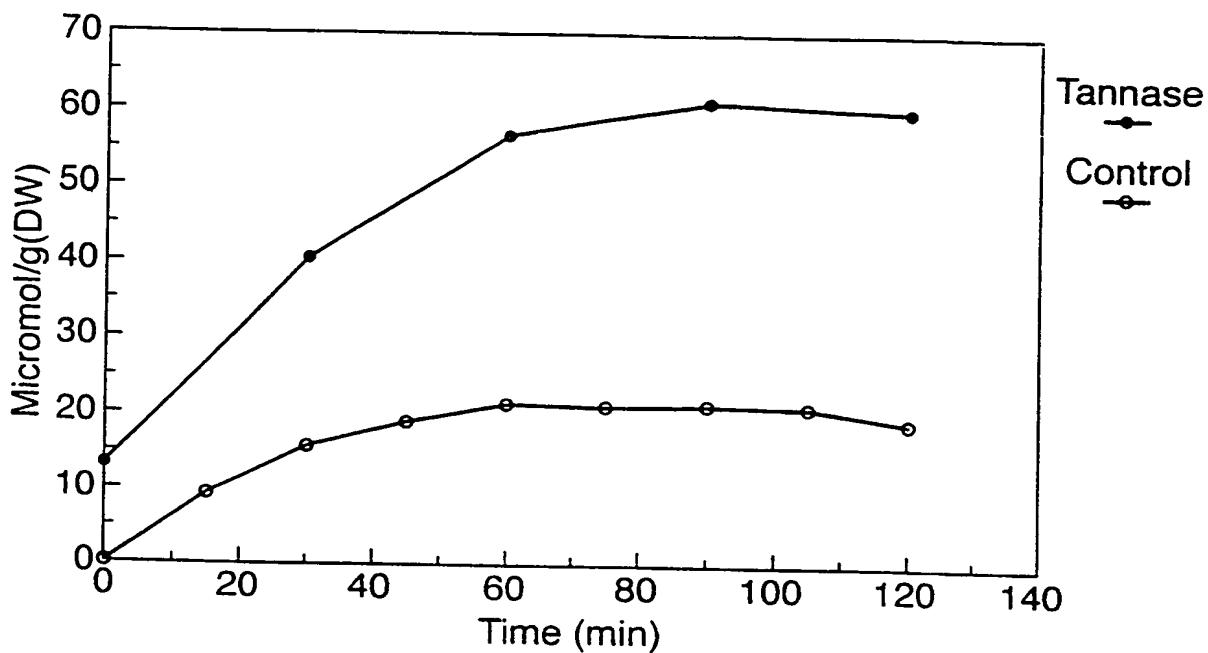


Fig.6.

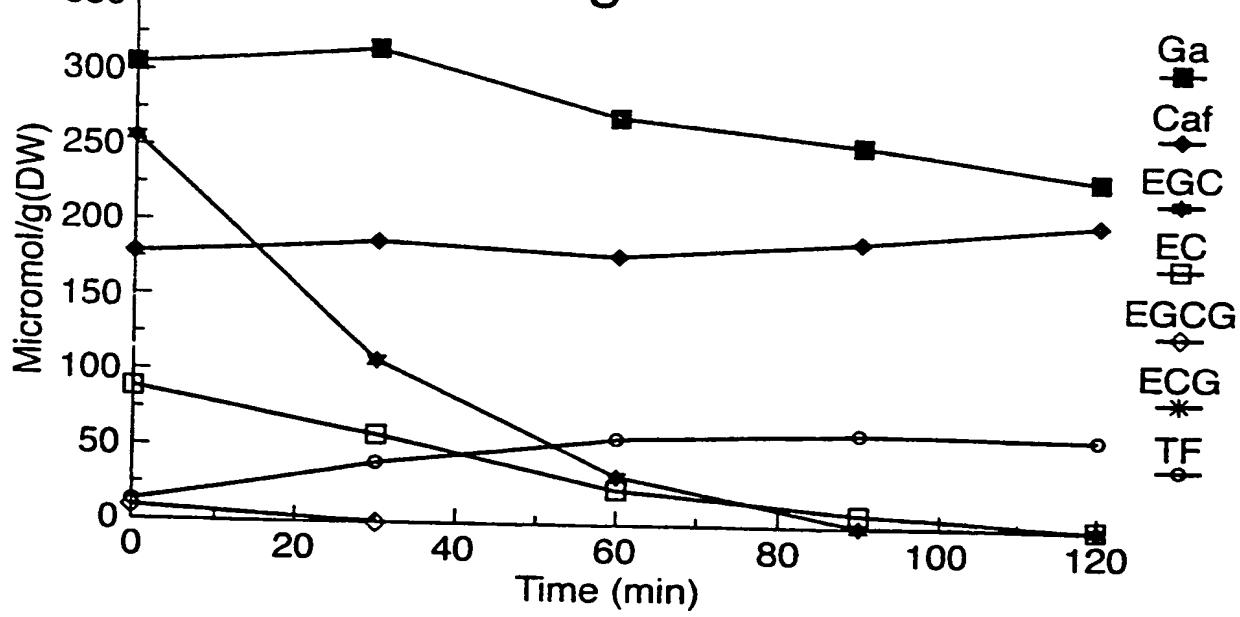


Fig.7.

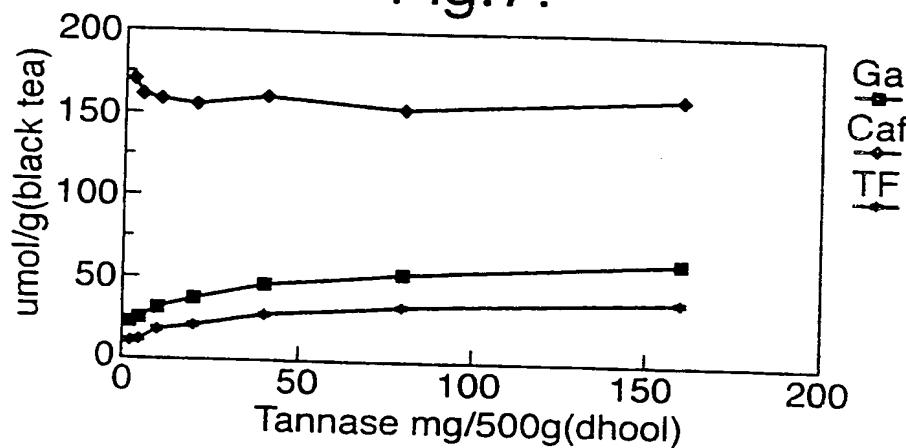


Fig.8.

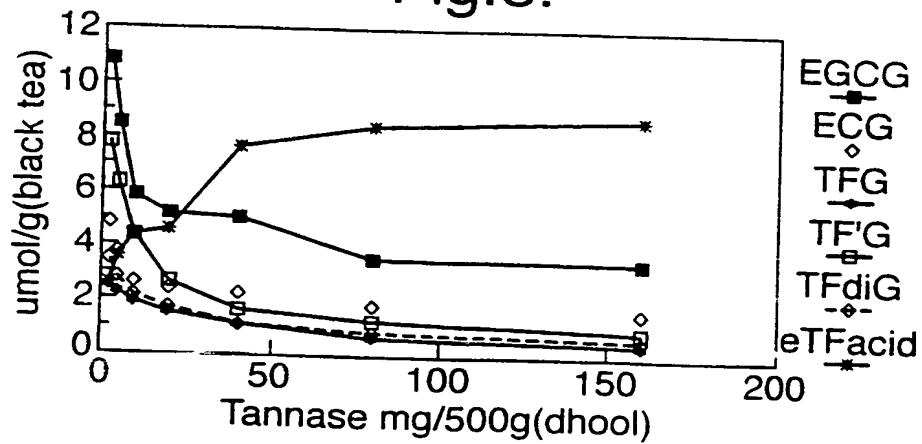


Fig.9a.

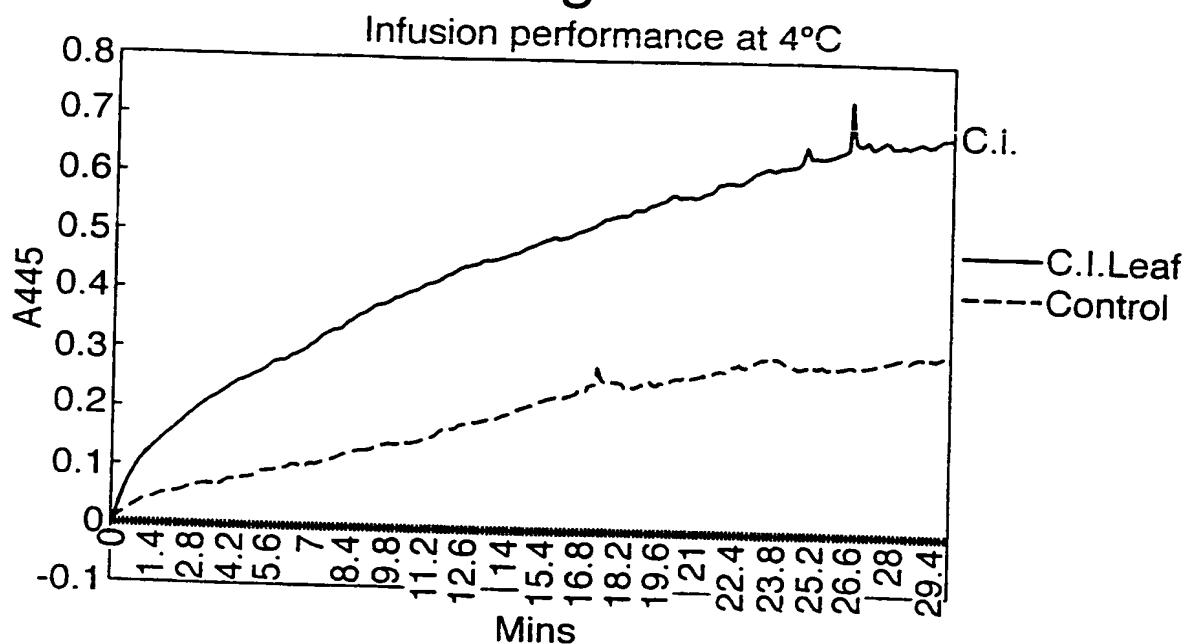
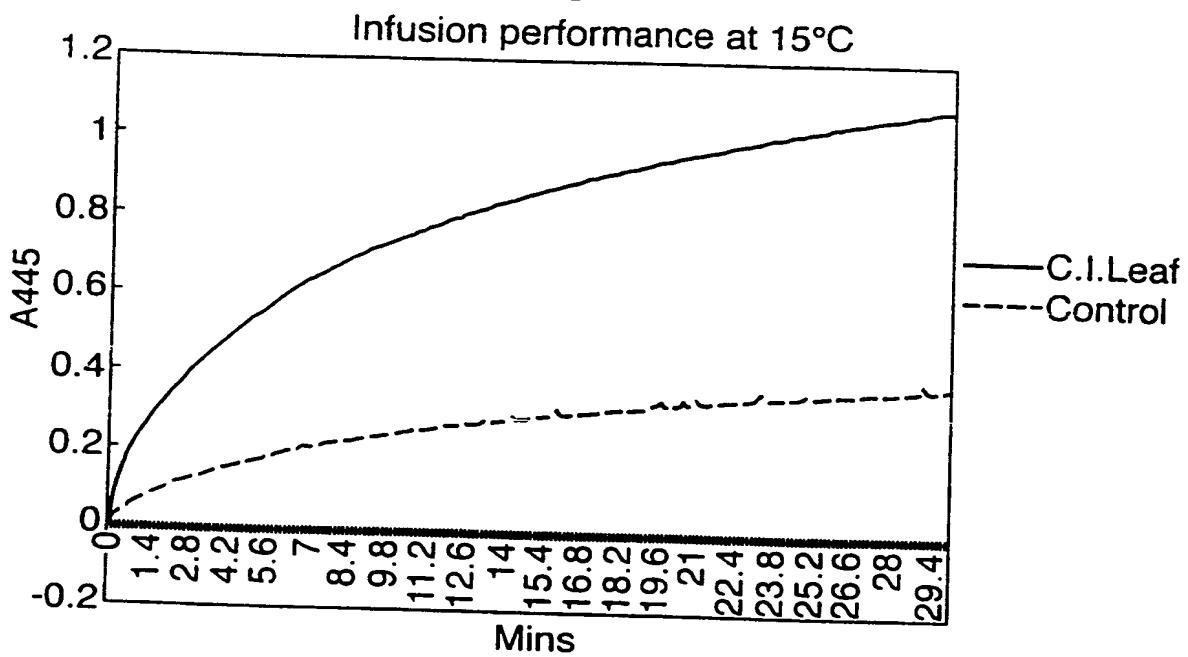


Fig.9b.



6/7

Fig.9c.

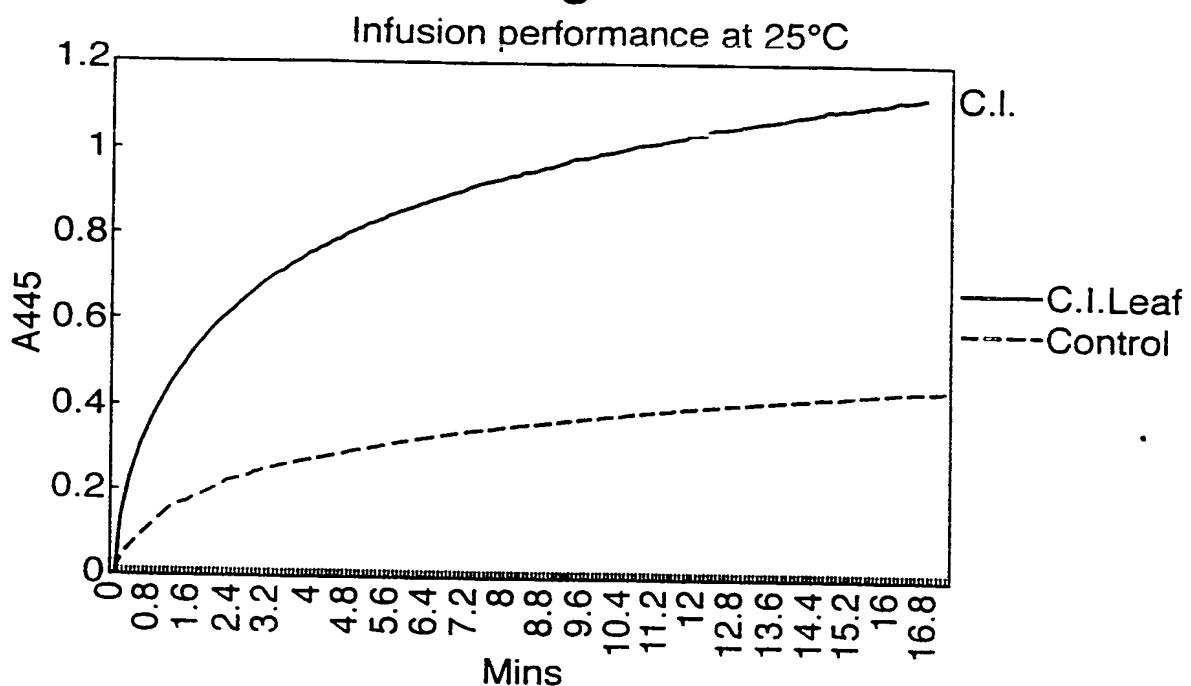


Fig.9d.

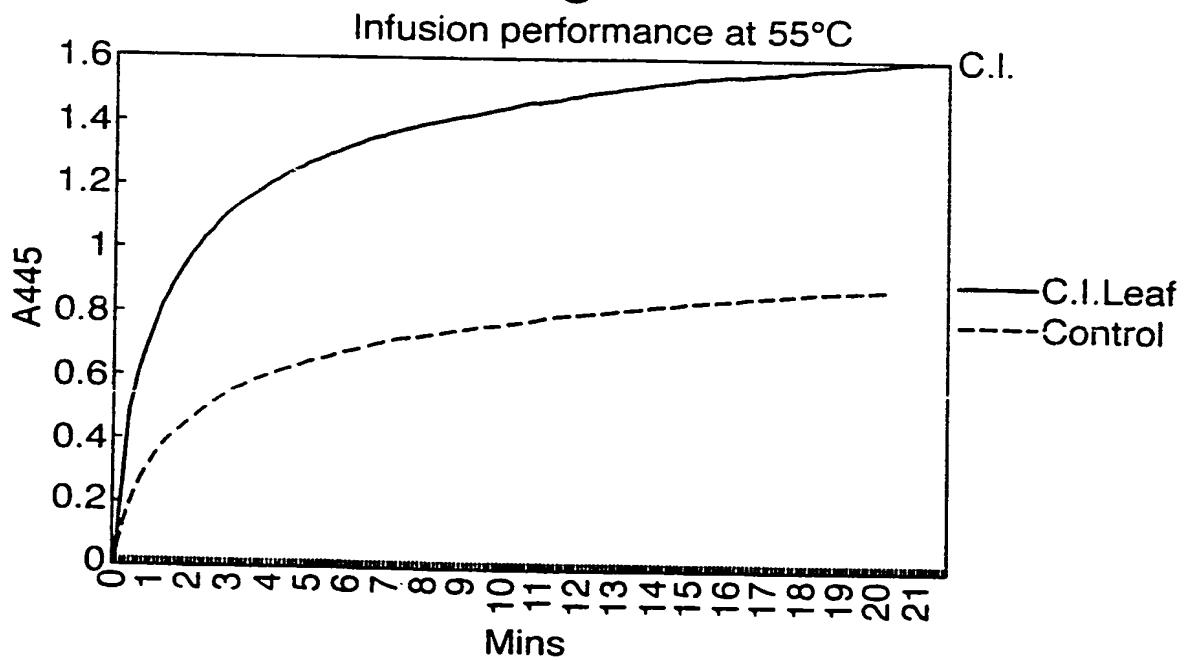
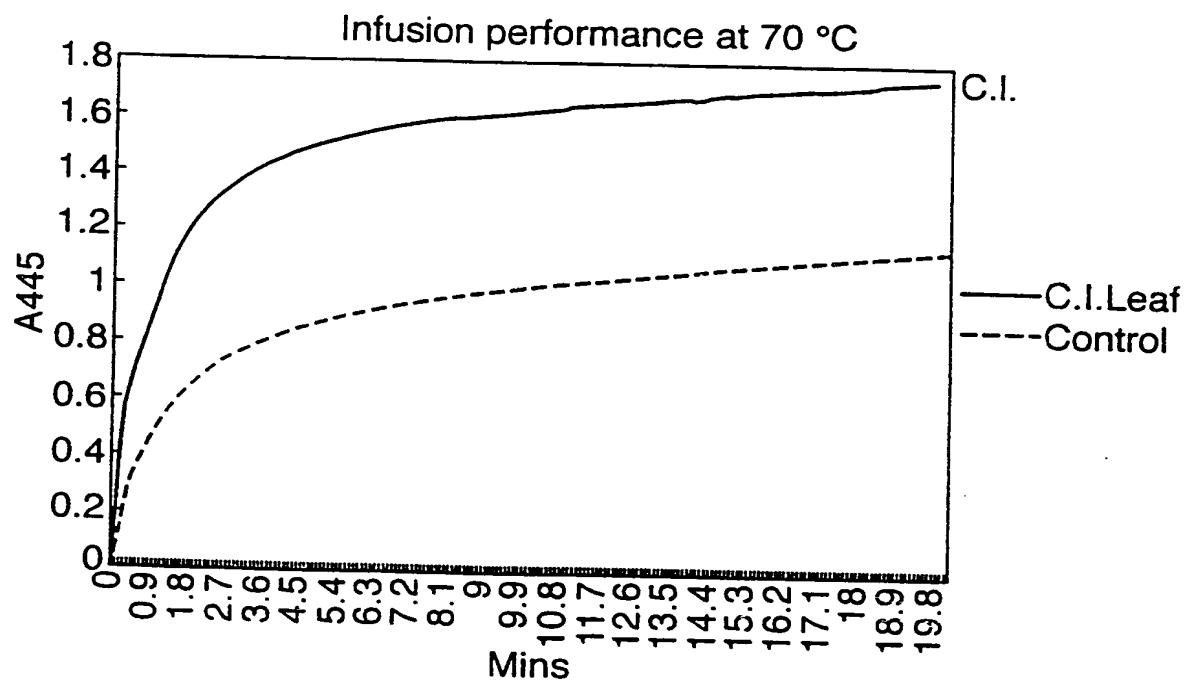


Fig.9e.



## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/GB 00/00359

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 7 A23F3/10

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 A23F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 3 812 266 A (SANDERSON G ET AL) 21 May 1974 (1974-05-21) cited in the application claim 5; example VIII	1-3,5, 11,13
A	US 4 051 264 A (SANDERSON GARY WARNER ET AL) 27 September 1977 (1977-09-27) cited in the application claims 12,14	4,6-10, 12
A	US 3 903 306 A (FAIRLEY CHARLES JOHN ET AL) 2 September 1975 (1975-09-02) column 2, line 39 - line 61; claim 1	1-4,8, 11-13
		1,5-13
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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Date of the actual completion of the international search

9 May 2000

Date of mailing of the international search report

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## INTERNATIONAL SEARCH REPORT

International Application No

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## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 760 213 A (UNILEVER ) 5 March 1997 (1997-03-05) cited in the application abstract -----	1-4,8-10

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/GB 00/00359

Patent document cited in search report	Publication date	Patent family member(s)			Publication date
US 3812266	A 21-05-1974	AU 5149473	A	01-08-1974	
		CA 995512	A	24-08-1976	
		CH 570772	A	31-12-1975	
		DE 2304073	A	16-08-1973	
		GB 1413351	A	12-11-1975	
		JP 915043	C	21-07-1978	
		JP 48082098	A	02-11-1973	
		JP 52042877	B	27-10-1977	
		KE 2642	A	25-06-1976	
US 4051264	A 27-09-1977	AU 501876	B	05-07-1979	
		AU 1207976	A	22-09-1977	
		CA 1087446	A	14-10-1980	
		CH 622410	A	15-04-1981	
		DE 2610533	A	07-10-1976	
		GB 1546508	A	23-05-1979	
		IN 144499	A	13-05-1978	
		JP 51115999	A	13-10-1976	
US 3903306	A 02-09-1975	AU 452833	B	20-08-1974	
		AU 3969272	A	13-09-1973	
		CA 963718	A	04-03-1975	
		GB 1319745	A	06-06-1973	
		JP 54140797	A	01-11-1979	
		ZA 7201456	A	29-11-1972	
EP 0760213	A 05-03-1997	US 5879730	A	09-03-1999	